## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application.

## Listing of Claims:

## 1. (Currently Amended) A compound of formula I

$$R_4$$
  $X$   $(R_3)_n$   $NH_2$   $(I)$ ,  $R_1$   $N$   $N$ 

wherein

n is from 0 to 4,

R<sub>1</sub> is hydrogen, unsubstituted or substituted lower alkyl or halogen,

R<sub>2</sub> is in the 4 position of the cyclohexane ring and is hydroxy, amino, N,N-di-lower alkylamino, pyrimidinyl-amino, 1,4,5,6-tetrahydro-pyrimidinyl-amino, 4,5-dihydro-1H-imidazolyl-amino, azetidin-1-yl, pyrrolidin-1-yl, 1-piperidyl, lower alkyl-piperazin-1-yl, morpholin-4-yl, thiomorpholin-4-yl; a radical R<sub>5</sub>-(C=Y)-NH-, wherein R<sub>5</sub> is lower alkyl, lower alkoxy, amino, Nlower alkylamino, N-(phenyl-lower alkyl)-amino, N-(lower alkyl-phenyl-lower alkyl)-amino, N-(lower alkoxy-phenyl-lower alkyl)-amino, N-(morpholin-4-vl-lower alkyl)-amino, N-(N', N'-dilower alkylamino-lower alkyl)-amino, lower alkoxy-lower alkoxy, 1-piperidyl-lower alkyl, morpholin-4-yl-lower alkyl or lower alkyl-piperazin-1-yl-lower alkyl, and Y is oxygen or imino; or a radical R6-sulfonylamino, wherein R6 is lower alkyl or N,N-di-lower alkylamino, hydroxy; unsubstituted, mono or disubstituted amino; an optionally substituted heterocyclic radical containing at least one nitrogen ring atom and being attached to the cyclohexane ring of the molecule of formula I via a nitrogen ring atom; a radical Rs-(C=Y)-NH-, wherein Rs is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino, a heterocyclic radical, or etherified hydroxy, and Y is oxygen, sulfur or imino; or a radical R6sulfonylamino, wherein Re is unsubstituted or substituted lower alkyl, unsubstituted, mono-or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro,

R<sub>3</sub> is lower alkyl, hydroxy, amino- or halogen substituted lower alkyl, hydroxy, cyano, lower alkoxy, lower alkanoyloxy, amino, mono- or di-lower alkylamino, lower alkanoylamino, carboxy, lower alkoxycarbonyl or halogen, wherein the R<sub>3</sub> substituents can be selected independently of one another if m 1;

R4 is benzyl, a-radical R2-CR8(R9), wherein R2 is cyclobutyl, cyclopentyl, cyclohexyl, phenyl, furyl, pyrrolyl, thienyl or pyridyl, said R2-substitutents being optionally substituted by one or more radicals selected from lower-alkyl and halogen, and R8 and R9 are independently of each other hydrogen, lower-alkyl or halogen, and

X is selected from -O-, -NH and S-, or a salt thereof.

2. (Cancelled) A compound of formula I according to claim 1, wherein n is from 0 to 4,

Rt-is-hydrogen, unsubstituted or substituted lower alkyl or halogen,

R<sub>2</sub>-is-hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical containing at least one nitrogen ring atom and being attached to the cyclohexane ring of the molecule of formula I via a nitrogen ring atom; a radical R<sub>5</sub> (C=Y) NH, wherein R<sub>5</sub> is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino, a heterocyclic radical, or etherified hydroxy, and Y is oxygen, sulfur or imino; or a radical R<sub>6</sub>-sulfonylamino, wherein R<sub>6</sub> is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro; R<sub>3</sub> is lower alkyl or lower alkoxy, wherein the R<sub>3</sub>-substituents can be selected independently of one another if n>1.

R<sub>4</sub> is a radical R<sub>2</sub>-GR<sub>8</sub>(R<sub>9</sub>)-, wherein R<sub>2</sub> is cyclobutyl, cyclopentyl, cyclohexyl, phenyl, furyl, pyrrolyl, thienyl, pyridyl or phenyl substituted by one or more radicals selected from lower alkyl and halogen, and R<sub>8</sub> and R<sub>9</sub> are independently of each other hydrogen, lower alkyl or halogen, and

X is selected from O, NH and S, or a salt thereof:

3. (Cancelled) A compound of formula I according to claim 1, wherein is 0, R<sub>1</sub>-is-hydrogen, unsubstituted or substituted lower alkyl or halogen, R<sub>2</sub>-is hydroxy; unsubstituted, mono- or disubstituted amino; an optionally substituted heterocyclic radical containing at least one nitrogen ring atom and being attached to the cyclohexane ring of the molecule of formula I via a nitrogen ring atom; a radical R<sub>5</sub>- (G=Y) NHI, wherein R<sub>5</sub>-is unsubstituted or substituted lower alkyl, unsubstituted, mono- or disubstituted amino, a heterocyclic radical, or etherified hydroxy, and Y is oxygen, sulfur or imino; or a radical R<sub>5</sub>-

sulfonylamino, wherein R<sub>6</sub> is unsubstituted or substituted lower alkyl, unsubstituted, mono-or disubstituted amino-or phenyl optionally substituted by lower alkyl, lower alkoxy-or nitro, R<sub>4</sub> is benzyl, and X is selected from O , NH- and S , or a salt-thereof.

4. (Cancelled) A compound of formula I according to claim 1, wherein n is 0,

R<sub>1</sub> is hydrogen, unsubstituted or substituted lower-alkyl or halogen,

R<sub>2</sub> is hydroxy; unsubstituted, mono—or disubstituted amino; an optionally substituted heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical to the cyclohexane ring of the molecule of formula I occurs via a nitrogen ring atom; a radical R<sub>5</sub> (G=Y) NIH, wherein R<sub>5</sub> is lower alkyl, unsubstituted, mono—or disubstituted amino, etherified hydroxy, a heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical occurs via a nitrogen ring atom, lower alkyl substituted by said heterocyclic radical or by one or more radicals selected independently of one another from the group consisting of amino, N lower alkylamino, N,N dilower alkylamino, N lower alkanoylamino, N,N dilower alkanoylamino, hydroxy, lower alkoxy, lower alkoxy, lower alkoxy, lower alkanoyl, lower alkanoyloxy, cyano, nitro, carboxy, lower alkoxycarbonyl, carbamoyl, amidino, guanidino, ureido, mercapto, lower alkylthio and halogen, and Y is oxygen, sulfur or imino; or a radical R<sub>6</sub>-sulfonylamino, wherein R<sub>6</sub>-is unsubstituted or substituted lower alkyl, unsubstituted, mono—or disubstituted amino or phenyl optionally substituted by lower alkyl, lower alkoxy or nitro,

R<sub>4</sub> is benzyl, and X is selected from O, NH and S, or a salt thereof.

5. (Cancelled) A compound of formula I according to claim-1, wherein n is 0,

R1-is-hydrogen, lower alkyl or halogen,

 $R_2$  is hydroxy; unsubstituted, mono—or disubstituted amino; an optionally substituted heterocyclic radical having from 4 to 8 ring members and from 1 to 3 heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical to the cyclohexane ring of the molecule of formula I occurs via a nitrogen ring atom; a radical  $R_3$  (C=Y) NH, wherein  $R_3$  is lower alkyl, unsubstituted or monosubstituted amino, etherified hydroxy, or lower alkyl substituted by a heterocyclic radical having from 4 to 8 ring members and from 1 to 3

heteroatoms whereby at least one heteroatom is nitrogen and the binding of the heterocyclic radical occurs via a nitrogen ring atom, and Y is oxygen or imino; or a radical R<sub>6</sub>-sulfonylamino, wherein R<sub>6</sub>-is lower-alkyl or disubstituted amino,

R4-is benzyl, and

X is selected from O, NH and S, or a salt thereof.

6. (Cancelled) A compound of formula I according to claim 1, wherein n is 0,

R+ is hydrogen, lower alkyl or halogen,

R<sub>2</sub> is hydroxy, amino, N,N di lower alkylamino, pyrimidinyl amino, 1,4,5,6 tetrahydro pyrimidinyl amino, 4,5 dihydro 1H imidazolyl amino, azetidin 1-yl, pyrrolidin 1-yl, 1-piperidyl, lower alkyl-piperazin 1-yl, morpholin 4-yl, thiomorpholin 4-yl, a radical R<sub>5</sub> (C=Y) NH, wherein R<sub>5</sub> is lower alkyl, lower alkoxy, amino, N lower alkylamino, N (phenyl lower alkyl) amino, N (lower alkyl phenyl lower alkyl) amino, N (lower alkoxy phenyl lower alkyl) amino, N (morpholin 4-yl lower alkyl) amino, N (N',N' di lower alkylamino-lower alkyl) amino, lower alkoxy lower alkoxy, 1-piperidyl lower alkyl, morpholin 4-yl lower alkyl or lower alkyl-piperazin 1-yl lower alkyl, and Y is oxygen or imino; or a radical R<sub>5</sub> sulfonylamino, wherein R<sub>5</sub> is lower alkyl or N,N di lower alkylamino,

R4-is benzyl, and

Xis O,

or a salt-thereof.

7. (Original) A compound of formula I according to claim 1, selected from the group consisting of cis-4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;

trans-4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol; cis-5-(3-benzyloxy-phenyl)-7-(4-piperidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-5-(3-benzyloxy-phenyl)-7-(4-piperidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-(4-pyrrolidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-pyrrolidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-[4-(4-methyl-piperazin-1-yl)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

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trans-5-(3-benzyloxy-phenyl)-7-[4-(4-methyl-piperazin-1-yl)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;
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cis-5-(3-benzyloxy-phenyl)-7-(4-morpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-morpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-7-(4-azetidin-1-yl-cyclohexyl)-5-(3-benzyloxy-phenyl)-7l-f-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-7-(4-azetidin-1-yl-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-(4-thiomorpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-thiomorpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-7-(4-diethylamino-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-7-(4-amino-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-7-(4-amino-cyclohexyl)-5-(3-benzyloxy-phenyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; cis-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-carbamic acid methyl ester;

cis-1-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-methylurea:

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-2-piperidin-1-yl-acetamide;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-2-morpholin-4-yl-acetamide;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-2-(4-methyl-piperazin-1-yl)-acetamide;

cis-5-(3-benzyloxy-phenyl)-7-[4-(pyrimidin-2-ylamino)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-[4-(1,4,5,6-tetrahydro-pyrimidin-2-ylamino)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-5-(3-benzyloxy-phenyl)-7-[4-(4,5-dihydro-1H-imidazol-2-ylamino)-cyclohexyl]-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-methanesulfonamide;

cis-N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-N,N-dimethylaminosulfonamide;

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cis-5-(3-benzyloxy-phenyl)-7-(4-dimethylamino-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-
ylamine;
N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-acetamide;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-ethyl-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohcxyl}-3-isopropyl-
urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-propyl-
urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo|2,3-d|pyrimidin-7-yl]-cyclohexyl}-3-butyl-urea;
cis-1-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(3-methyl-
benzyl)-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-benzyl-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(4-
methoxy-benzyl)-urea;
cis-1-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-tert-butyl-
urea;
cis- N-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-guanidine;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(2-
dimethylamino-ethyl)-urea;
cis-1-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(2-
morpholin-4-yl-ethyl)-urea;
cis-1-[4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-3-(3-
morpholin-4-yl-propyl)-urea;
cis-{4-[4-amino-5-(3-benzyloxy-phenyl)-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexyl}-carbamic acid
2-methoxy-ethyl ester;
cis-4-[4-amino-5-(3-benzyloxy-phenyl)-6-bromo-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
trans-4-[4-amino-5-(3-benzyloxy-phenyl)-6-bromo-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
cis-4-[4-amino-5-(3-benzyloxy-phenyl)-6-methyl-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
trans-4-[4-amino-5-(3-benzyloxy-phenyl)-6-methyl-pyrrolo[2,3-d]pyrimidin-7-yl]-cyclohexanol;
trans-5-(3-benzyloxy-phenyl)-6-methyl-7-[4-(4-methyl-piperazin-1-yl)-cyclohexyl]-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
trans-5-(3-benzyloxy-phenyl)-7-(4-dimethylamino-cyclohexyl)-6-methyl-7H-pyrrolo[2,3-
d]pyrimidin-4-ylamine;
trans-5-(3-benzyloxy-phenyl)-7-(4-diethylamino-cyclohexyl)-6-methyl-7H-pyrrolo[2,3-
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trans-5-(3-benzyloxy-phenyl)-6-methyl-7-(4-pyrrolidin-1-yl-cyclohexyl)-7H-pyrrolo[2,3-

d]pyrimidin-4-ylamine;

d]pyrimidin-4-ylamine;

trans-5-(3-benzyloxy-phenyl)-6-methyl-7-(4-morpholin-4-yl-cyclohexyl)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; trans-7-(4-azetidin-1-yl-cyclohexyl)-5-(3-benzyloxy-phenyl)-6-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-ylamine; and pharmaceutically acceptable salts thereof.

- 8. (Cancelled) A compound of formula I, or a pharmaceutically acceptable salt thereof, according to claim 1 for use in a method for the treatment of the human or animal body.
- 9. (Previously Presented) A pharmaceutical composition comprising a compound of formula I or a pharmaceutically acceptable salt thereof according to claim 1, together with at least one pharmaceutically acceptable carrier.

10.-12. (Cancelled)